

Sheet 1 of 1

<b>INFORMATION DISCLOSURE CITATION</b>	ATTY. DOCKET NO. <u>4662-88</u>	SERIAL NO. <u>10/554,294</u>
	APPLICANT	
	<u>MINK et al</u>	
(Use several sheets if necessary)	FILING DATE <u>Unassigned</u>	TC/A.U. <u>Unassigned</u>

## **U.S. PATENT DOCUMENTS**

## **FOREIGN PATENT DOCUMENTS**

**OTHER DOCUMENTS (including Author, Title, Date, Pertinent pages, etc.)**

	International Search Report
/BD/	Kelvin L. BAUMANN et al; "The Convergent Synthesis of CI-981, an Optically Active, Highly Potent, Tissue Selective Inhibitor of HMG-CoA Reductase"; <i>Tetrahedron Letters</i> , Vol. 33, No. 17, pp. 2283-2284; 1992
/BD/	Peter W.K. WOO et al; "Atorvastatin, An HMG-COA Reductase Inhibitor and Effective Lipid-Regulating Agent – Part III <sup>1a,b</sup> Syntheses of [ <sup>2</sup> H <sub>5</sub> ]-,[ <sup>13</sup> C <sub>B</sub> ], and [ <sup>13</sup> C <sub>7</sub> , <sup>15</sup> N] Atorvastatin and Their Application in Metabolic and Pharmacokinetic Studies; <i>Journal of Labelled Compounds and Radiopharmaceuticals J. Labelled Cpd. Radiopharm.</i> 42, 135-145; (1999)
/BD/	Philip L. BROWER et al; "The Synthesis of (4R-cis)-1,1-Dimethylethyl 6-cyanomethyl-2,2-dimethyl-1,3-dioxane-4-acetate, a Key Intermediate for the Preparation of CI-981, a Highly Potent, Tissue Selective Inhibitor of HMG-CoA Reductase"; <i>Tetrahedron Letters</i> , Vol. 33, No. 17, pp. 2279-1182; 1992
/BD/	William A. GREENBERG et al; "Development of an Efficient, Scalable, Aldolase-Catalyzed Process for Enantioselective Synthesis of Statin Intermediates"; <i>PNAS</i> , April 20, 2004; Vol. 101, No. 16, pp. 5788-5793

/Bernard Dentz/

06/02/2008

### Date Considered

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